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DATE: Wednesday, September 20, 2006

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<input type="checkbox"/>	L15	L14 and l1	1
<input type="checkbox"/>	L14	(548/335.1)! [CCLS]	511
<input type="checkbox"/>	L13	(435/6)! [CCLS]	36324
<input type="checkbox"/>	L12	L11 and l1	10
<input type="checkbox"/>	L11	L10 and l5	18
<input type="checkbox"/>	L10	apoptosis or cell death	46411
<input type="checkbox"/>	L9	L8 and l1	14
<input type="checkbox"/>	L8	imidazolyl disulfid\$	24
<input type="checkbox"/>	L7	imidazolyl	39407
<input type="checkbox"/>	L6	L5 and L1	10
<input type="checkbox"/>	L5	powis.in.	78
<input type="checkbox"/>	L4	L3 not @ay>1995	5
<input type="checkbox"/>	L3	L2 and antibod\$	97
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<input type="checkbox"/>	L1	thioredoxin	6843

END OF SEARCH HISTORY

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NEWS 9 JUN 02 The first reclassification of IPC codes now complete in INPADOC
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=> s thioredoxin
5812 THIOREDOXIN
3679 THIOREDOXINS
L1 6388 THIOREDOXIN
(THIOREDOXIN OR THIOREDOXINS)

=> s cancer? or tumor? or neoplas?
306312 CANCER?
441863 TUMOR?
463561 NEOPLAS?
L2 732005 CANCER? OR TUMOR? OR NEOPLAS?

=> s 11 (1) 12
L3 589 L1 (L) L2

=> s 13 not py>1995
10304536 PY>1995
L4 49 L3 NOT PY>1995

=> s imidazolyl?
L5 10560 IMIDAZOLYL?

=> s 15 and 14
L6 1 L5 AND L4

=> d ibib

L6 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER: 1995:523489 CAPLUS
DOCUMENT NUMBER: 123:47459
TITLE: The thioredoxin/thioredoxin reductase redox system and control of cell growth
AUTHOR(S): Powis, Garth; Oblong, John E.; Gasdaska, Pamela Y.; Berggren, Margareta; Hill, Simon R.; Kirkpatrick, D. Lynn
CORPORATE SOURCE: Arizona Cancer Center, Tucson, AZ, 85724, USA
SOURCE: Oncology Research (1994), 6(10-11), 539-44

PUBLISHER: CODEN: ONREE8; ISSN: 0965-0407
Elsevier
DOCUMENT TYPE: Journal
LANGUAGE: English

=> d ibib kwic

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AB Thioredoxin is a redox protein that is important for a variety
of intracellular functions, possibly including regulation of transcription
factor activity. We have shown that human thioredoxin has the
same predicted amino acid sequence as adult T-cell-derived leukemic cell
growth factor. Recombinant human thioredoxin stimulates the
proliferation of Swiss murine 3T3 fibroblasts with an EC50 of 100 nM and
the proliferation of a number of human cancer cells. Site-directed
mutagenesis of the active-site cysteines of thioredoxin has
shown that redox activity is necessary for the stimulation of cell
proliferation. Added 125I-thioredoxin is taken up by cells in
culture and could have intracellular action. A series of alkyl 2-
imidazolyl disulfides have been shown to be competitive inhibitors
of human thioredoxin reductase with Ki values of 3.3 to 8.6
μM. The compds. inhibited Swiss 3T3 serum-dependent proliferation with
IC50 values of 2.0 to 4.0 μM, and one compound inhibited Swiss 3T3
thioredoxin-dependent proliferation with an IC50 value of 60 nM.

IT Cell proliferation
Neoplasm
Redox reaction
(thioredoxin/thioredoxin reductase redox system and
control of cell growth)